

Comparative study of the synovial histology in rheumatoid arthritis, spondyloarthropathy, and osteoarthritis: influence of disease duration and activity

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Abstract

Objectives—To compare the macroscopic and microscopic characteristics of synovial tissue in rheumatoid arthritis (RA), spondyloarthropathy (SpA), and osteoarthritis (OA) after exclusion of possible biases induced by disease duration or activity, or both.

Methods—Synovial biopsy specimens were obtained by needle arthroscopy in patients with early RA (n=16), late RA (n=14), early SpA (n=23), and OA (n=12). Macroscopic and microscopic features were scored on a four point scale and analysed as a function of disease duration (early versus late RA), local and systemic disease activity, and diagnosis.

Results—Except for the maximal synovial lining thickness, no significant differences were seen between early and late RA. For disease activity, synovial histology was only weakly correlated with C reactive protein in RA, but seemed to be strongly dependent on effusion of the biopsied joint in all disease groups. After stratification for local disease activity, no disease related differences were found in patients without joint effusion. In contrast, important differences were found between patients with RA and SpA with active joint effusion. Synovial vascularity was macroscopically increased in SpA versus RA ($p=0.017$). A straight vessel pattern was only seen in RA, while tortuous vessels were preferentially seen in SpA. Vascularity was also microscopically increased in SpA compared with RA ($p=0.031$), and correlated with the macroscopic vascularity ($r_s=0.36$, $p=0.036$). CD3+ ($p=0.008$), CD4+ ($p=0.008$), and CD20+ ($p=0.024$) lymphocytes were overrepresented in RA compared with SpA. The integrin expression in RA was characterised by a decrease of $\alpha V\beta 3$ in the synovial lining ($p=0.006$) and an increase of $\alpha V\beta 5$ in the sublining ($p<0.001$).

Conclusions—The immune architecture of the synovial membrane is more dependent on local disease activity than on disease duration. Synovium obtained from clinically affected joints shows important histological differences between RA and SpA.

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Synovitis is a major characteristic of chronic inflammatory joint diseases of autoimmune origin, such as rheumatoid arthritis (RA) and spondyloarthropathy (SpA). It can also occur as a secondary inflammatory symptom in osteoarthritis (OA), which is primarily induced by biomechanical stress on cartilage and subchondral bone. Studies in RA indicate that the synovial membrane has a dominant role in the joint inflammation and destruction, as suggested by the changes in synovial histology: (a) thickening of the synovial lining layer, as a result of infiltration by CD68+ cells¹ and both proliferation and reduced apoptosis of type B synoviocytes²⁻⁵; (b) neovascularisation of the sublining layer⁶⁻⁸; (c) infiltration of the sublining with T and B lymphocytes,⁹⁻¹² plasma cells,^{10,12} and macrophages^{1,12}; and (d) alteration of the adhesion molecule expression, including the expression of αV integrins which may have a role in both neovascularisation and pannus formation.^{13,14}

These observations suggest that the synovial membrane is both the primary site of inflammation, triggered by autoreactive T cells and macrophages, and the main effector organ, as the hyperplastic "aggressive" pannus leads to cartilage and bone erosion.¹⁵⁻¹⁷ However, most of these histological studies have used synovial tissue obtained at the time of surgery, thereby creating a selection bias of the patient group for disease duration and activity.^{18,19} It remains, therefore, unclear if the observed histological changes are of primary pathogenic relevance, or if they are secondary bystander phenomena of the chronic joint inflammation and destruction. Moreover, the use of surgery specimens also biased the choice of the control group towards OA, and most studies failed to include other inflammatory arthritides, such as SpA. Thus both the specificity of the observations in RA and the histology of the synovial membrane in SpA remain largely to be investigated.

Owing to the development of needle arthroscopy, which made it possible to visualise and sample the synovial membrane in early and late stages of all types of peripheral joint diseases,²⁰ this study aims at describing disease related histological features of the synovial membrane. Firstly, it investigates the influence of disease duration (early versus late RA) and activity (presence of joint effusion, sedimentation rate, C reactive protein (CRP), swollen joint count) on synovial histology. Secondly, after correction for these possible biases by appropriate

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stratification it compares systematically the synovial lining hypertrophy, the vascularity, the cellular infiltration, and the αV integrin expression in RA, SpA, and OA.

Materials and methods

PATIENTS

Synovial tissue samples were obtained from 71 consecutive patients undergoing needle arthroscopy of the knee for pain or synovitis, or both. Thirty patients were diagnosed as RA, according to the American College of Rheumatology (ACR) criteria.²¹ Of these patients with RA, 16 had early disease, defined as a disease duration of less than one year as measured from the first symptoms of arthritis, and 14 had a longstanding disease (>1 year, range 1.5–20). Twenty three patients were diagnosed as SpA, according to the European Spondyloarthropathy Study Group criteria²²: five patients with ankylosing spondylitis, 10 with psoriatic arthritis, and eight undifferentiated spondyloarthropathies. All patients with SpA had a disease duration of less than one year. OA was diagnosed in 12 patients, according to the ACR criteria.²³ The remaining six patients were excluded from the study as no definite diagnosis could be made. Age, sex, disease duration, and use of drugs were recorded. The total number of swollen joints (including hands, wrists, elbows, shoulders, knees, ankles, and feet) and the presence of joint effusion in the investigated knee were evaluated clinically: effusion in the investigated knee was considered to be a primary variable for local disease activity as other symptoms of synovial inflammation (warmth, swelling,...) were virtually always associated with the presence of synovial fluid. Laboratory investigations included the measurement of serum CRP, erythrocyte sedimentation rate, and rheumatoid factor, and HLA typing. Joint destruction was assessed by *x* ray, using the Steinbrocker criteria.²⁴ Table 1 summarises the features of the different patient groups.

Synovial biopsy specimens were obtained by needle arthroscopy as described previously.²⁰ Briefly, needle arthroscopy of the knee was performed under local anaesthesia using a 2.7 mm Hopkins rod-lens telescope (Karl Storz, Tuttlingen, Germany). The joint cavity was carefully inspected, and synovial vascularity

and hypertrophy were scored on a four point semiquantitative scale by an experienced arthroscopist (DB or FVdB).²⁰ The morphology of the blood vessels was described as a straight, tortuous, or mixed pattern.²⁵ Synovial membrane biopsy specimens (n=10) were obtained from the macroscopically inflamed areas with a 2.7 mm biopsy forceps (Karl Storz). In each patient, five biopsy specimens were stored in formaldehyde and embedded in paraffin. Sections, 5 μ m, were cut and stained with haematoxylin-eosin for histological analysis. The remaining five biopsy specimens were snap frozen and mounted in Jung tissue freezing medium (Leica Instruments, Nussloch, Germany). Frozen sections were cut and mounted on Star Frost glass slides (Knittelgläser, Braunschweig, Germany). Both paraffin embedded and frozen sections were used for immunohistochemistry.

IMMUNOHISTOCHEMISTRY

Paraffin embedded sections were dewaxed, heated in a microwave oven in citrate buffer (0.1 mol/l, pH 6.0) for antigen retrieval, and incubated for 60 minutes with the following mouse monoclonal antibodies (mAbs): anti-CD20 (clone L26, Dako, Glostrup, Denmark), anti-plasma cell (clone VS38c, Dako), anti-CD68 (clone PG-M1, Dako). Frozen sections were fixed for 10 minutes in acetone and incubated for 30 minutes with the following mouse mAbs: anti-CD3 (clone UCHT1, Dako), anti-CD4 (clone MT310, Dako), anti-CD8 (clone DK25, Dako), anti- $\alpha V\beta 3$ (clone 23C6, Pharmingen, San Diego, USA), and anti- $\alpha V\beta 5$ (clone P1F6, Chemicon International Inc, Temecula, USA). Parallel sections were incubated with irrelevant isotype matched mAb as negative control. After rinsing, the endogenous peroxidase was blocked with 1% hydrogen peroxide. The sections were subsequently incubated for 15 minutes with a biotinylated antimouse secondary antibody, followed by 15 minutes with a streptavidin-peroxidase complex (LSAB+ Kit, Dako). The colour reaction was developed with 3-amino-9-ethylcarbazole substrate (Dako) as chromogen. Finally, the sections were counterstained with haematoxylin. All incubations were carried out at room temperature and the sections were washed with phosphate buffered saline between all steps.

Table 1 Clinical data of the patients with early rheumatoid arthritis (early RA; disease duration of less than one year), late rheumatoid arthritis (late RA; disease duration of more than one year), spondyloarthropathy (SpA), and osteoarthritis (OA). Mean (standard deviation) (range)

	Early RA	Late RA	SpA	OA
Number of patients	16	14	23	12
Sex (M/F)	6/10	3/11	13/10	5/7
Age (years)	51 (17) (18–80)	56 (12) (36–81)	41 (15) (19–64)	62 (15) (38–80)
Disease duration (years)	0.6 (0.3) (0.2–1)	8.5 (5.7) (1.5–20)	0.7 (0.5) (0.1–1)	Unknown
Joint effusion of the knee (+/–)	10/6	8/6	13/10	3/9
Number of swollen joints	6.9 (4.7)	6.8 (6.6)	1.6 (0.8)	1.2 (0.9)
Serum CRP* (mg/l)	36 (26)	42 (41)	24 (23)	05 (03)
Sedimentation rate (mm/1st h)	35 (25)	38 (18)	24 (18)	13 (9)
Shared epitope (+/–)	12/4	11/3	6/17	3/9
HLA-B27 (+/–)	0/16	1/13	11/12	0/12
Rheumatoid factor (+/–)	6/10	10/4	0/23	0/12
<i>x</i> Ray score	0.80 (0.86)	1.17 (1.11)	0.23 (0.69)	1.33 (1.07)
NSAID* (+/–)	13/3	6/8	12/11	4/8
DMARD* (+/–)	1/15	6/8	8/15	0/12
Corticosteroids (+/–)	5/11	6/8	2/21	0/12

*CRP = C reactive protein; NSAID = non-steroidal anti-inflammatory drug; DMARD = disease modifying antirheumatic drug.

MICROSCOPIC ANALYSIS

Stained sections were coded and analysed by two independent observers who were unaware of the diagnosis and clinical data. The analysis included all areas of the five biopsy specimens and a global score was given for each variable. A semiquantitative four point scale was developed according to previously used scoring systems for synovial tissue^{10 12 26-28}; zero represented the lowest and three the highest level of expression. As some histological markers are more abundant than others in synovial tissue, the scoring system was calibrated for each marker separately by examining a representative number of synovial samples. Thus the sensitivity of the scoring system was different for each marker, allowing comparison of a specific marker between the different patient groups, but not comparison of different markers in one group. This semiquantitative scale was used for all variables, unless indicated otherwise. The synovial lining layer was scored for the number of CD68+ cells, the mean lining layer thickness, and the maximal lining layer thickness. Mean and maximal lining thickness were scored separately as it appeared that synovial lining layer hyperplasia can be focal and thus that both variables can be discrepant in some samples. The mean thickness of the synovial lining layer was evaluated by counting the number of cell layers in six randomly selected regions and calculating the mean.²⁹ The maximal thickness was scored semiquantitatively (0:1-2; 1:3-4; 2:5-6; 3:≥7 cell layers).^{10 27} In the sublining the vascularity was evaluated by the number of blood vessels and their localisation (superficial, profound, diffuse). Infiltration of the sublining layer with inflammatory cells was evaluated by the global number of infiltrating cells, the number of lymphocytes, plasma cells, and neutrophils, and the presence of lymphoid aggregates (+/-). Additionally, immunohistochemical stainings were scored for CD68, CD20, CD3, CD4, CD8, and plasma cells. Finally, the expression of the integrins α V β 3 and α V β 5 was scored in three compartments: synovial lining layer, sublining layer, and

endothelium. The scores obtained by the two observers were concordant in more than 95% of cases. When discordant scores were obtained, which differed by a maximum of one point, the mean of the two scores was used.

STRATIFICATION

The following factors that might influence the comparison of RA, SpA, and OA samples were analysed in the first part of the study: disease duration, presence of effusion in the biopsied joint, CRP, sedimentation rate, and swollen joint count. Factors that seemed significantly to influence the histological features were subsequently used for stratification of the second part of the study in order to avoid systematic biases in the comparison of RA, SpA, and OA.

STATISTICAL ANALYSIS

To avoid statistical biases created by multiple comparisons, the Kruskal-Wallis test was used for comparing the mean scores in the different patient groups; only when this test showed significant differences, were the exact p values calculated with the Mann-Whitney U test and corrected for the number of comparisons. Spearman's test was used for correlations. $p < 0.05$ was considered to be significant.

Results

INFLUENCE OF DISEASE DURATION ON SYNOVIAL HISTOLOGY IN RA

To determine whether the macroscopic and microscopic features of the synovial membrane were dependent on disease duration, the 16 patients with early RA were compared with the 14 patients with late RA. No significant differences in macroscopic vascularity or villous hypertrophy were seen. At the microscopic level the only difference was an increased maximal lining thickness in late RA (mean (SEM) 2.50 (0.19)) compared with early RA (1.73 (0.20), $p = 0.013$), whereas the mean lining thickness was not increased in late (2.26 (0.34)) *v* early RA (2.30 (0.33)). As no other significant differences were seen, either in vas-

Table 2 Microscopic features of the synovial membrane in rheumatoid arthritis (RA) with or without joint effusion, spondyloarthropathy (SpA) with or without joint effusion, and osteoarthritis (OA) with or without joint effusion. Mean histological score (standard error of the mean)

	RA with joint effusion (n=18)	RA without joint effusion (n=12)	SpA with joint effusion (n=13)	SpA without joint effusion (n=10)	OA with joint effusion (n=3)	OA without joint effusion (n=9)
Mean lining thickness	2.69 (0.30)*	1.68 (0.31)*	3.48 (0.66)	2.02 (0.35)	2.47 (0.79)	1.40 (0.18)
Max lining thickness	2.31 (0.18)	1.71 (0.29)	1.36 (0.20)	1.33 (0.33)	1.67 (0.33)	1.25 (0.25)
CD68 in lining	1.06 (0.24)	0.77 (0.26)	1.69 (0.24)*	0.81 (0.27)*	1.33 (0.67)	0.89 (0.26)
α V β 3 in lining	1.09 (0.31)†	1.20 (0.58)	2.40 (0.16)†	2.25 (0.25)	1.33 (0.33)	1.00 (0.41)
α V β 5 in lining	2.73 (0.14)	2.50 (0.50)	2.55 (0.16)	1.80 (0.49)	2.67 (0.33)	1.5 (0.87)
Vascularity	1.58 (0.14)*†	1.04 (0.18)*	2.23 (0.29)†	1.60 (0.27)	2.00 (0.00)	0.78 (0.22)
Infiltration	1.92 (0.15)*	0.59 (0.18)*	2.08 (0.25)*	0.80 (0.20)*	1.50 (0.29)	0.22 (0.15)
Neutrophils	0.66 (0.17)	0.27 (0.20)	1.31 (0.35)	0.45 (0.24)	0.67 (0.33)	0.11 (0.11)
Lymphocytes	1.92 (0.09)*	0.86 (0.23)*	1.88 (0.26)*	0.95 (0.26)*	1.67 (0.33)	0.22 (0.15)
Plasma cells	0.90 (0.23)*	0.09 (0.09)*	0.38 (0.15)	0.00 (0.00)	0.00 (0.00)	0.11 (0.11)
CD68 in sublining	1.34 (0.28)	0.68 (0.27)	1.23 (0.20)*	0.28 (0.15)*	0.67 (0.33)	0.33 (0.24)
CD38	0.44 (0.19)	0.17 (0.17)	0.56 (0.31)	0.14 (0.14)	1.07 (0.73)	0.33 (0.33)
CD20	0.94 (0.27)†	0.14 (0.10)	0.15 (0.15)†	0.22 (0.15)*	0.00 (0.00)	0.11 (0.11)
CD3	2.41 (0.16)*†	0.64 (0.24)*	1.55 (0.27)*†	0.58 (0.27)*	1.00 (0.00)	0.72 (0.28)
CD4	1.91 (0.19)*†	0.32 (0.19)*	0.95 (0.24)†	0.25 (0.17)	1.00 (0.58)	0.44 (0.18)
CD8	1.62 (0.20)*	0.45 (0.16)*	1.40 (0.28)*	0.08 (0.08)*	1.00 (0.58)	0.44 (0.18)
α V β 5 in sublining	2.45 (0.21)*†	0.50 (0.50)*	0.55 (0.25)†	0.00 (0.00)	0.33 (0.00)	0.00 (0.00)
α V β 3 on endothelium	1.64 (0.34)	0.4 (0.4)	1.20 (0.33)	1.00 (0.71)	0.00 (0.00)	0.00 (0.00)
α V β 5 on endothelium	0.46 (0.25)	0.0 (0.0)	0.09 (0.09)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)

* $p < 0.05$ for comparison of patients with and without joint effusion within one diagnostic group. † $p < 0.05$ for comparison of patients with RA with joint effusion and patients with SpA with joint effusion.

cularity, inflammatory infiltration, or αV integrin expression, the patients with early and late RA were considered as one group for further analysis.

INFLUENCE OF LOCAL DISEASE ACTIVITY ON SYNOVIAL HISTOLOGY

As synovial membrane samples were obtained both from patients with active joint effusion and patients with only knee pain, which might be due to secondary cartilage damage rather than to synovial inflammation, the influence of local disease activity on synovial histology was

analysed by comparing patients with and without effusion of the biopsied joint. Table 2 summarises these results. In RA, most histological variables were increased in the 18 patients with joint effusion compared with the 12 patients without joint effusion, reaching significance for mean lining thickness ($p=0.040$), vascularity ($p=0.023$), and inflammatory infiltration ($p<0.001$), number of lymphocytes ($p<0.001$), plasma cells ($p=0.013$), CD3+ cells ($p<0.001$), CD4+ cells ($p<0.001$), and CD8+ cells ($p=0.001$), and expression of $\alpha V\beta 5$ in the sublining layer ($p=0.027$) (fig 1A). Lymphoid

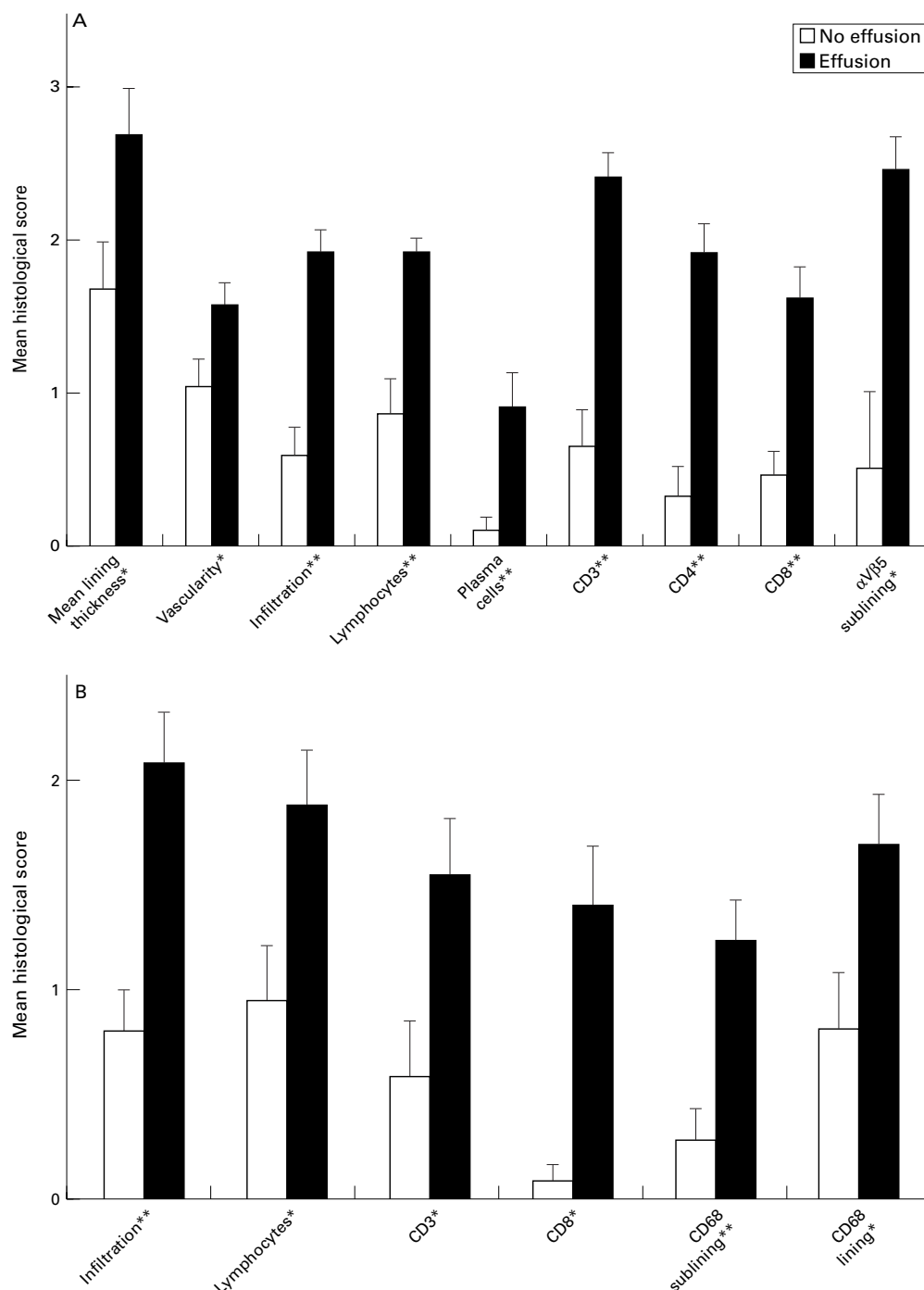


Figure 1 Comparison of the histological scores in patients with and without effusion of the biopsied joint. Mean (standard error of the mean). * $p<0.05$. ** $p<0.01$. (A) Patients with rheumatoid arthritis with ($n=18$) and without joint effusion ($n=12$). (B) Patients with spondyloarthropathy with ($n=13$) and without joint effusion ($n=10$).

Table 3 Macroscopic morphology of the synovial blood vessels in rheumatoid arthritis (RA) and spondyloarthropathy (SpA): straight vessels, tortuous vessels, or mixed pattern. Results are shown as number (%)

	RA (n=18)	SpA (n=13)
Straight	5 (28)	0 (0)
Tortuous	5 (28)	8 (62)
Mixed	8 (44)	5 (38)

aggregates were seen in eight patients with RA with joint effusion (44%), but in none of the patients without effusion. A similar trend was found in the SpA group, with an increase in degree of infiltration ($p=0.003$), number of lymphocytes ($p=0.030$), CD3+ cells ($p=0.031$), CD8+ cells ($p=0.011$), CD68+ cells in the lining layer ($p=0.025$), and CD68+ cells in the sublining layer ($p=0.006$) (fig 1B). Also in OA strong vascularity and infiltration was only found in patients with joint effusion, but these differences did not reach significance owing to the small number of observations (only three patients with joint effusion). Because of the significant impact of local disease activity on the synovial histology in all patient groups, further comparisons were stratified for the presence or absence of effusion in the biopsied joint.

INFLUENCE OF SYSTEMIC DISEASE ACTIVITY ON SYNOVIAL HISTOLOGY

Systemic disease activity, as evaluated by CRP serum concentrations, sedimentation rate, and number of swollen joints, had less influence on the histological findings. In RA, the CRP level correlated with the degree of inflammatory infiltration ($r_s=0.46$, $p=0.013$), the number of lymphocytes ($r_s=0.41$, $p=0.027$), CD3+ cells ($r_s=0.44$, $p=0.021$), CD4+ cells ($r_s=0.58$,

$p=0.002$), CD8+ cells ($r_s=0.54$, $p=0.003$), CD68+ cells in the sublining layer ($r_s=0.52$, $p=0.007$), $\alpha V\beta 3$ expression on endothelium ($r_s=0.77$, $p=0.001$), and $\alpha V\beta 5$ expression in the sublining layer ($r_s=0.71$, $p=0.006$). Similar correlations were found with the sedimentation rate, but not with the number of swollen joints. However, in SpA and OA there were no significant correlations between systemic variables of disease activity and histological findings. Moreover, when this analysis was corrected for the influence of local disease activity, by including only patients with active effusion of the biopsied joint, the only significant observations were the correlation in patients with RA of CRP with the number of CD4+ cells ($r_s=0.54$, $p=0.031$), CD8+ cells ($r_s=0.50$, $p=0.049$), and $\alpha V\beta 3$ expression on endothelium ($r_s=0.64$, $p=0.033$).

MACROSCOPIC AND MICROSCOPIC FEATURES OF THE SYNOVIAL MEMBRANE IN RA VERSUS SPA, WITHOUT EFFUSION OF THE BIOPSIED JOINT

When the patients without active joint effusion were analysed, no differences were noted in macroscopic or microscopic features of the synovial membrane between RA ($n=12$), SpA ($n=10$), and OA ($n=9$) (table 2).

MACROSCOPIC FEATURES OF THE SYNOVIAL MEMBRANE IN PATIENTS WITH RA VERSUS PATIENTS WITH SPA WITH EFFUSION OF THE BIOPSIED JOINT

OA was not included in this analysis owing to the small sample number of patients with active joint effusion ($n=3$). Patients with RA ($n=18$) and SpA ($n=13$) with clinical joint effusion showed a similar degree of villous hypertrophy (1.31 (0.26) *v* 1.38 (0.29)), but the macro-

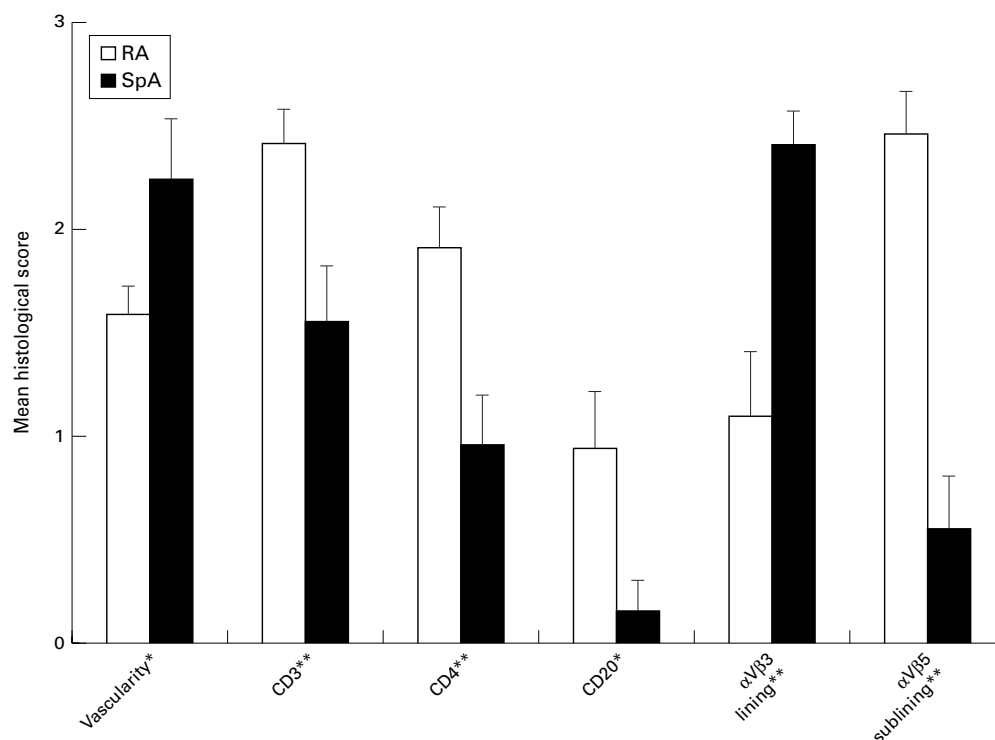


Figure 2 Comparison of the histological scores in 18 patients with rheumatoid arthritis (RA) and 13 patients with spondyloarthropathy (SpA) with effusion of the biopsied joint. Mean (standard error of the mean). * $p<0.05$. ** $p<0.01$.

scopic vascular pattern was clearly different. The vascularity was significantly increased in SpA (2.08 (0.28)) compared with RA (1.22 (0.19), $p=0.017$). Moreover, the morphology of the blood vessels was different: straight vessels were preferentially seen in RA, whereas tortuous vessels were predominant in SpA. However, a manifest tortuous pattern was also seen in 5/18 (28%) of the patients with RA, and a mixed pattern was common in both groups (table 3).

MICROSCOPIC FEATURES OF THE SYNOVIAL MEMBRANE IN PATIENTS WITH RA VERSUS PATIENTS WITH SPA WITH EFFUSION OF THE BIOPSIED JOINT

Table 2 shows the histological and immunohistochemical data and fig 2 illustrates the main differences between RA and SpA. No differences were observed in the synovial lining layer: mean lining thickness, maximal lining thickness, and number of CD68+ cells were similar in both groups. In the sublining layer,

vascularity was increased in SpA (2.23 (0.29)) compared with RA (1.58 (0.14), $p=0.031$), thus confirming the macroscopic observations. Microscopic and macroscopic vascularity correlated weakly ($r_s=0.36$, $p=0.036$). Both in RA and SpA, the blood vessels were preferentially seen in the superficial layers just beneath the synovial lining layer. The overall degree of inflammatory infiltration of the sublining layer was similar in RA and SpA. There were no significant differences in number of macrophages (CD68+), plasma cells, and neutrophils. However, RA synovium showed an important lymphocytic infiltration with an increase in number of CD3+ cells (2.41 (0.16) *v* 1.55 (0.27), $p=0.008$), CD4+ cells (1.91 (0.19) *v* 0.95 (0.24), $p=0.008$), and CD20+ cells (0.94 (0.27) *v* 0.15 (0.15), $p=0.024$). The number of CD8+ cells was not increased. Lymphoid aggregates were seen in 8/18 patients with RA, but in only 2/13 patients with SpA. Finally, the integrins $\alpha V\beta 3$ and $\alpha V\beta 5$ were differentially

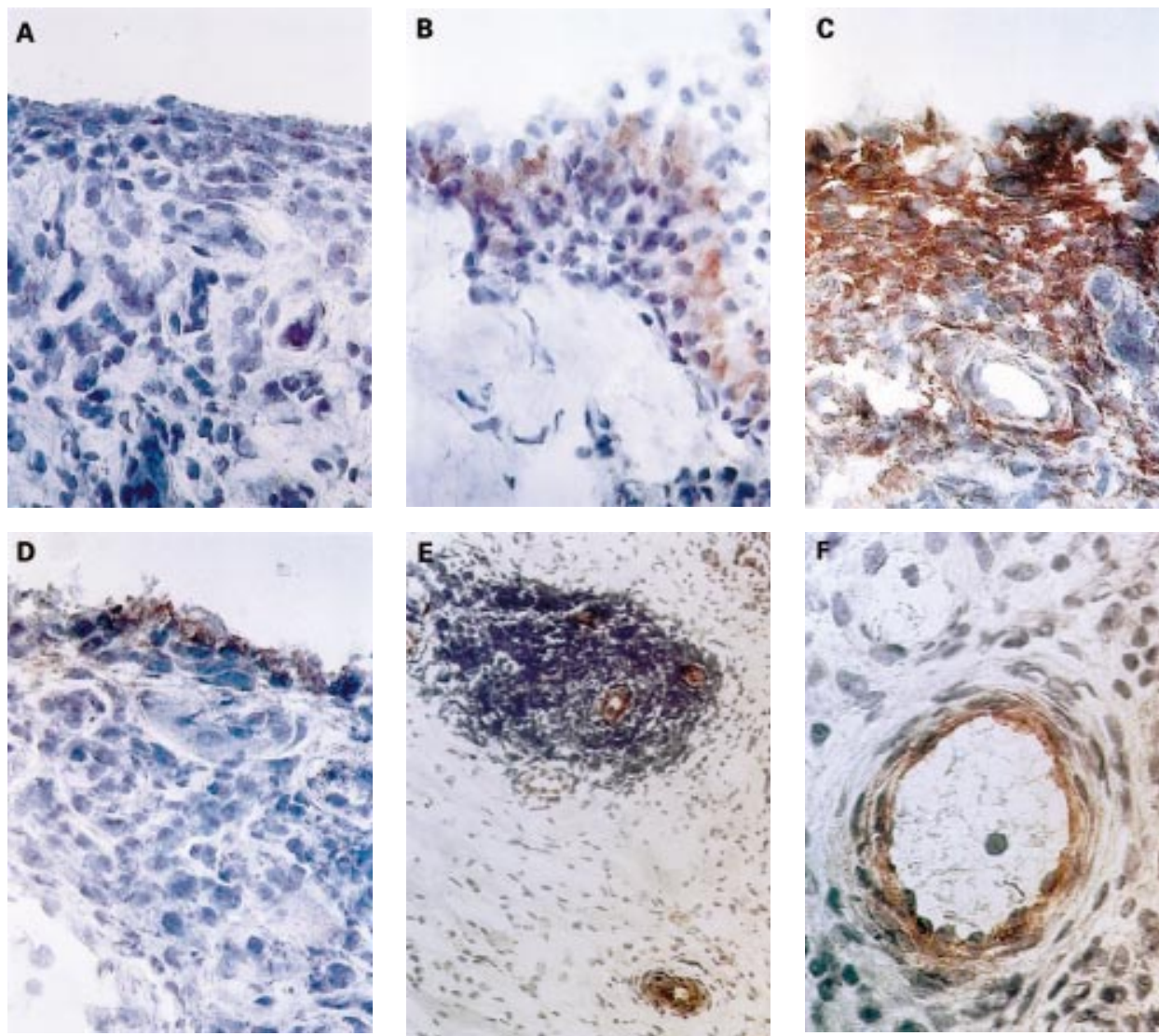


Figure 3 αV integrin expression in the synovial membrane. Frozen sections of synovial biopsy specimens from RA (A and C) and SpA (B and D) were stained immunohistochemically for $\alpha V\beta 3$ (A and B) and $\alpha V\beta 5$ (C and D). (A) $\alpha V\beta 3$ expression in RA synovium: no staining of synovial lining cells. (B) $\alpha V\beta 3$ expression in SpA synovium: staining of synovial lining cells. (C) $\alpha V\beta 5$ expression in RA synovium: staining of superficial synovial lining cells and of sublining cells. (D) $\alpha V\beta 5$ expression in SpA synovium: staining of superficial synovial lining cells, but not of sublining cells. (E) $\alpha V\beta 3$ expression on endothelial cells in RA synovium. (F) $\alpha V\beta 3$ expression on endothelial cells in SpA synovium.

expressed in both patient groups (fig 3). In the synovial lining layer, the expression of $\alpha V\beta 3$ was decreased in RA (1.09 (0.31)) *v* SpA (2.40 (0.16), $p=0.006$). In the sublining layer, the expression of $\alpha V\beta 5$ was increased in RA (2.45 (0.21)) *v* SpA (0.55 (0.25), $p<0.001$). The expression of both integrins on endothelium of sublining blood vessels was similar in both groups.

Discussion

Although the synovial membrane in RA has been analysed in numerous studies, little is known about the pathogenic relevance and specificity of the histological findings, such as synovial lining layer hyperplasia, neovascularisation, and inflammatory infiltration of the sublining with lymphocytes and macrophages. Either the use of surgical specimens created a possible bias with regard to disease duration and activity, or the studies failed to include an inflammatory, immune mediated joint disease as control. Our study was designed to exclude such biases by the use of needle arthroscopic biopsy specimens, and to describe subsequently disease-specific histological changes of the synovial membrane which might be of pathogenic or diagnostic relevance in RA and SpA.

A systematic comparison of early RA, defined as disease duration of less than one year, with established RA showed no major macroscopic or microscopic differences. The only exception, the increase of maximal lining layer thickness with disease duration, should be interpreted carefully as it might be explained by chance owing to multiple testing. These data are concordant with previous reports on inflammatory infiltration, T cell activation, and expression of adhesion molecules and cytokines in early and late RA,^{10 12 28 30} and support the concept that a number of histological and immunohistochemical findings are independent of disease duration. However, the present histological data do not allow us to conclude that the earliest phase of RA, which probably precedes the first clinical manifestations, and established disease are driven by similar immunological mechanisms.^{31 32}

To analyse the influence of local disease activity on synovial histology, patients with knee joint effusion and patients with knee pain only (without effusion) were studied. A first observation is that synovial tissue from knees without effusion can already depict some histological changes, such as inflammatory infiltration and increased vascularity, suggesting that the pain is not only due to secondary cartilage damage but also to synovial inflammation. This is not unexpected as histological changes were reported in clinically unaffected joints of patients with RA,^{8 33 34} suggesting an early pathogenic role for CD68+ macrophages and macrophage-derived proinflammatory cytokines in RA synovitis. Secondly, the inflammatory infiltration, as well as other histological variables, is strongly increased in patients with RA, SpA, and OA with joint effusion compared with patients without effusion. These data support the view that synovial his-

tology is influenced by local disease activity,¹² and question the use of non-inflammatory controls in comparative histology of the synovial membrane.

After stratification for local disease activity, systemic measures of disease activity, such as CRP and sedimentation rate, correlated only weakly with some histological features in RA but not in SpA and OA, which might be explained by the fact that they are not good measures of disease activity in SpA and OA.³⁵

Beside disease duration and disease activity, drugs might also be a confounding factor in the histological study of synovial tissue as numerous previous studies have shown the effect of antirheumatic drugs on synovial histology.³⁶⁻⁴¹ Although this issue was not analysed here and needs certainly to be considered in future studies, the fact that the use of non-steroidal anti-inflammatory drugs, disease modifying antirheumatic drugs, and corticosteroids was comparable in the different patient groups makes it unlikely that a possible drugs bias accounts for important differences in the comparison of RA, SpA, and OA.

When stratifying for local disease activity, no significant differences were found between patients with RA, SpA, and OA without joint effusion, which suggests that either these histological changes are not disease-specific, or that the semiquantitative approach used in this study is not sufficiently sensitive to detect small differences. In contrast, several significant disease related features were detected in the patients with RA and SpA with active joint effusion. Although it has to be considered that macroscopic scoring of the synovial membrane has not yet been validated, analysis of the synovial membrane during needle arthroscopy confirmed our previous report of an increased vascularity in SpA compared with RA.²⁰ A recent report indicates that there is also a difference in morphology of the blood vessels between these two pathologies: tortuous vessels were seen in 73% of SpA and 11% of RA, whereas straight vessels were seen in 89% of RA and 26% of SpA.²⁵ Our study confirms this observation, despite some minor discrepancies: (a) a straight vessel pattern is quite specific for but not very common in RA; (b) tortuous vessels are predominant in SpA, but are also seen in RA; and (c) an important proportion of both groups has a mixed vascularity pattern. The increased vascularity in SpA compared with RA was confirmed at the microscopic level and is in agreement with previous studies in PsA⁴² and SA.⁴³ Taken together, the increased macroscopic and microscopic vascularity and the different morphology of the blood vessels indicate that neovascularisation might be mediated by different growth factors and/or receptors in RA and SpA, though the present study showed no difference in αV integrin expression on endothelium. Furthermore, the present data indicate that neovascularisation might be even more important as a pathogenic mechanism and possible therapeutic target in SpA than in RA.

Conflicting results have been reported for the inflammatory infiltration of the synovium

in RA compared with SpA.^{10 28 44 45} These discrepancies might be explained by the differences in SpA subtypes or the differences in microscopic evaluation, or both. This study indicates that manifest lymphocytic infiltration is present in both diseases, but that CD3+ cells, CD4+ cells, CD20+ cells, and lymphoid aggregates are clearly increased in RA. Interestingly, the score for CD8+ cells is not different. Although the study was not designed to compare the level of expression of two markers in the same tissue, this might reflect an increased CD4/CD8 ratio in RA. Beside lymphocytes, no differences are seen in other cell types infiltrating the sublining layer, but although the relatively small number of patients in each SpA subgroup allows only preliminary observations, a slight but significant increase of neutrophils was noted in PsA compared with other SpA subtypes and RA (data not shown).

Finally, the expression of the integrins $\alpha V\beta 3$ and $\alpha V\beta 5$ was compared between RA and SpA. These integrins can be expressed on endothelial cells, where they mediate angiogenesis,^{46 47} and on synovial fibroblasts.^{12 48} In this study both integrins are expressed to a similar degree on endothelium in RA and SpA, indicating an important synovial neovascularisation in both diseases. On synovial fibroblasts, however, RA is characterised by a strongly significant decrease of $\alpha V\beta 3$ in the synovial lining layer and increase of $\alpha V\beta 5$ in the sublining layer. As engagement of αV integrins regulates proliferation, migration, and collagenase expression of a variety of cell types,⁴⁸⁻⁵² this differential integrin expression may have an important role in the aggressive growth of the synovial pannus in RA.

In conclusion, synovial histology is dependent on local disease activity but not on disease duration in clinically manifest RA. Comparative histology after stratification for this confounding factor allows the description of disease related changes of the synovial membrane, including neovascularisation in SpA, T and B cell infiltration in RA, and differential αV integrin expression. The presence of a disease related immune architecture of the synovial membrane warrants further investigation of the pathophysiological and/or diagnostic relevance of these findings.

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